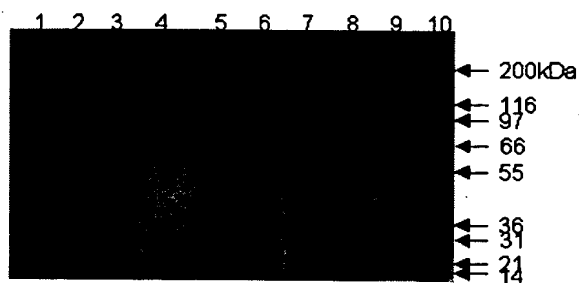
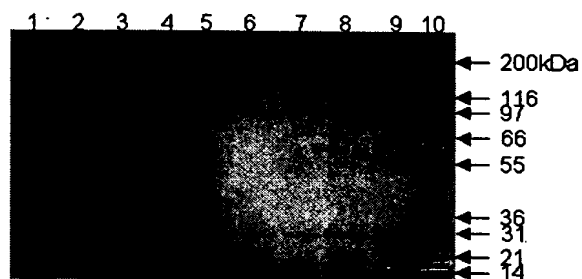


Fig. 1



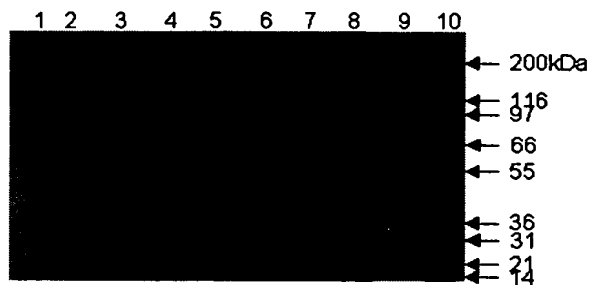
Lanes 1 & 10, marker proteins; lane 2 untreated mbh; lane 3, 50°C, lane 4, 60°C, lane 5, 70°C; lane 6, 80°C; lane 7, 90°C; lane 8, 100°C; lane 9, Protease M.

Fig. 2



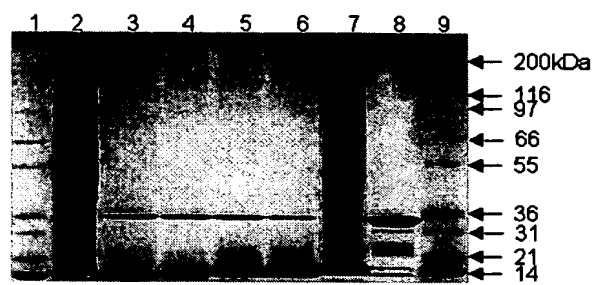
Lanes 1 & 10, marker proteins; lane 2 untreated mbh; lane 3, pH2, lane 4, pH4, lane 5, pH6; lane 6, pH8; lane 7, pH10; lane 8, pH12; lane 9, Protease M.

Fig. 3



Lanes 1 & 10, marker proteins;
lane 2 untreated mbh; lanes 3 - 8,
Rokko digest (20mg.ml^{-1} -
 0.1mg.ml^{-1}), lane 9, Rokko
(1mg.ml^{-1}).

Fig. 4



Lanes 1 & 9, marker proteins; lane 2
untreated mbh; lane 3, 2% SDS; lane
4, 1% SDS; lane 5, 0.5% SDS; lane 6,
0.25% SDS; lane 7, mbh + 2% SDS;
lane 8, Rokko (20mg.ml^{-1}).

Fig. 5

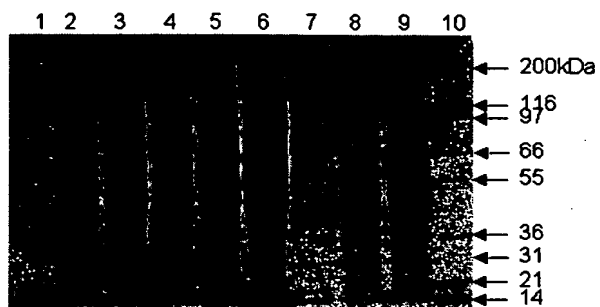
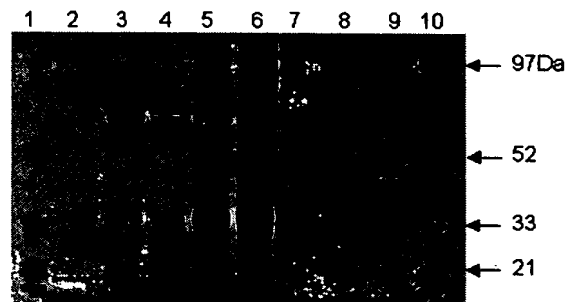


Fig. 6



Lanes 1 & 10, marker proteins; lanes 2 & 3, mbh; lanes 4 - 6, mbh pellet; lanes 7 - 9, mbh supernatant.

Fig.7

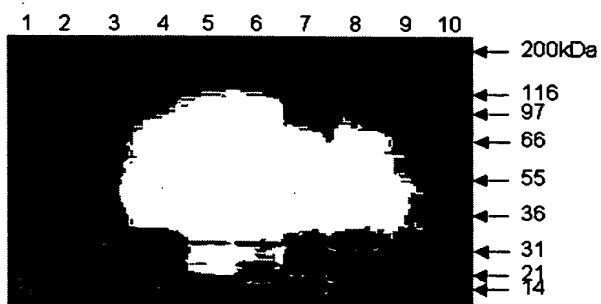


Fig.8



Lanes 1 & 10, marker proteins; lane 2, untreated mbh; lane 3, Protease G digest; lane 4, Protease G; lane 5, Protease R digest; lane 6, Protease R; lane 7, Protease C digest; lane 8, Protease C; lane 9, rec. mouse PrP.

1

2

3

4



Fig. 9

5

6

7

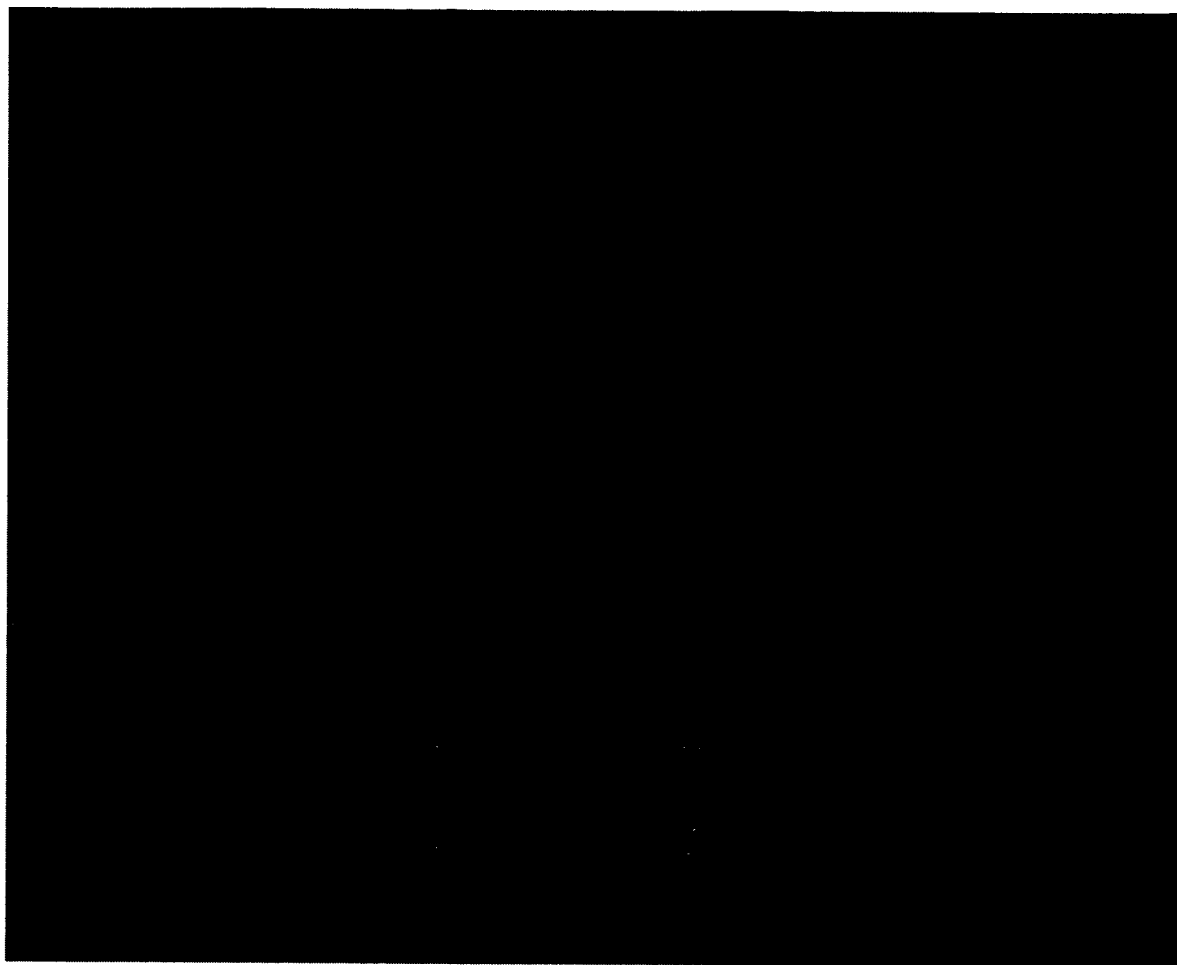


Fig. 10

1

2

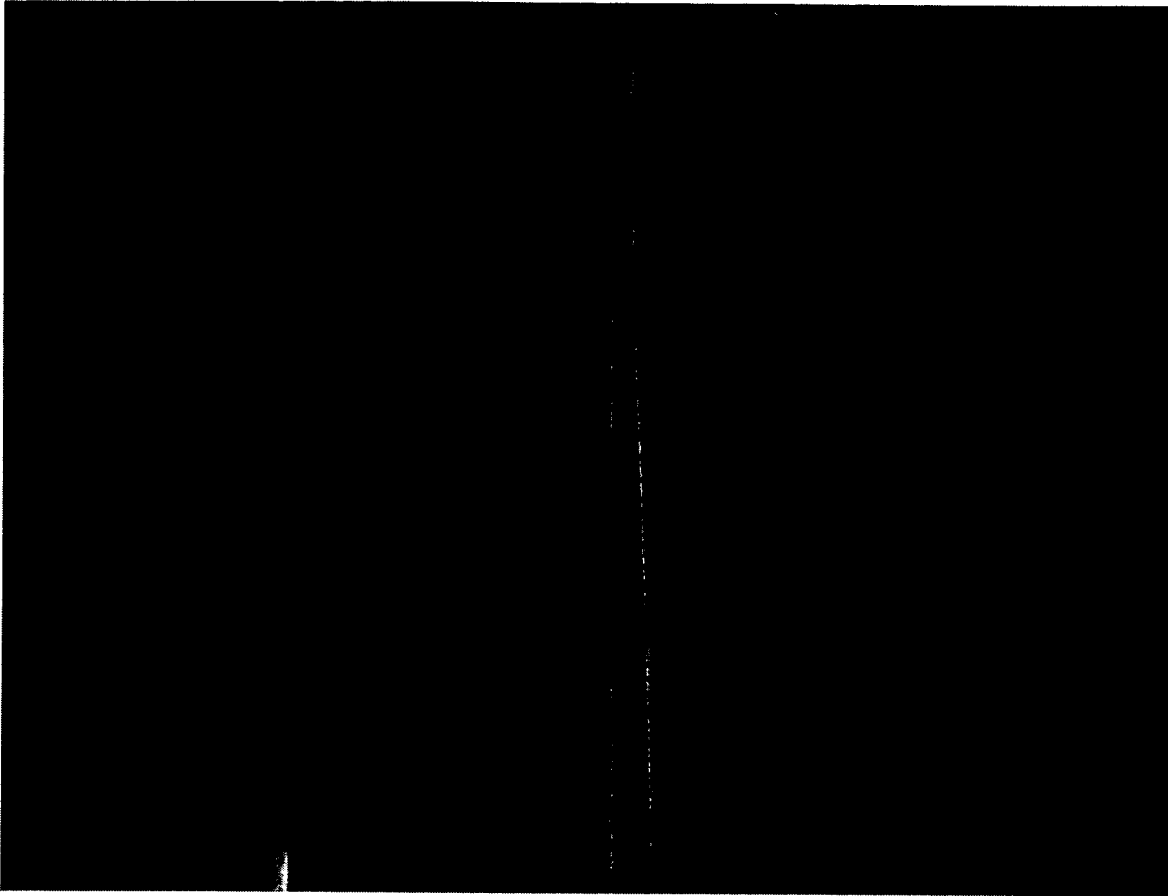


Fig. 11

3

7

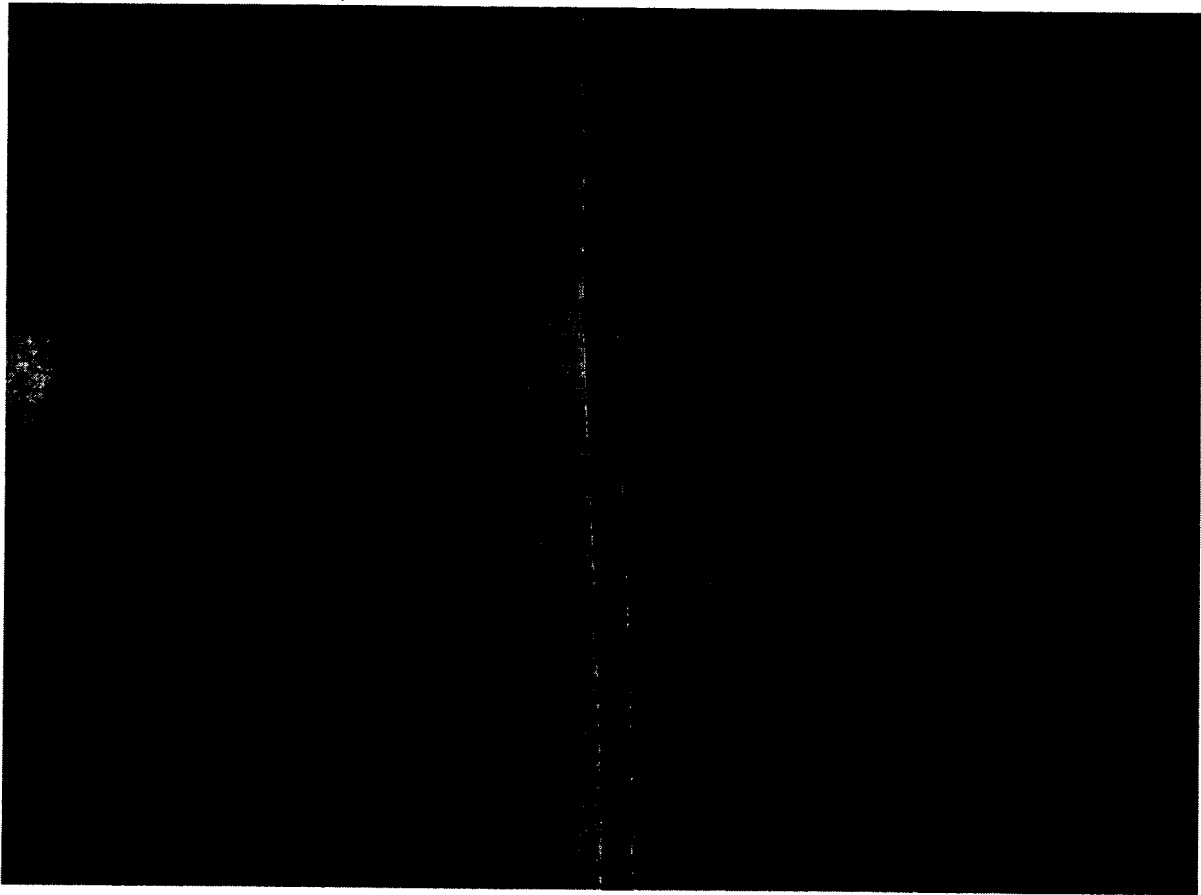


Fig. 12

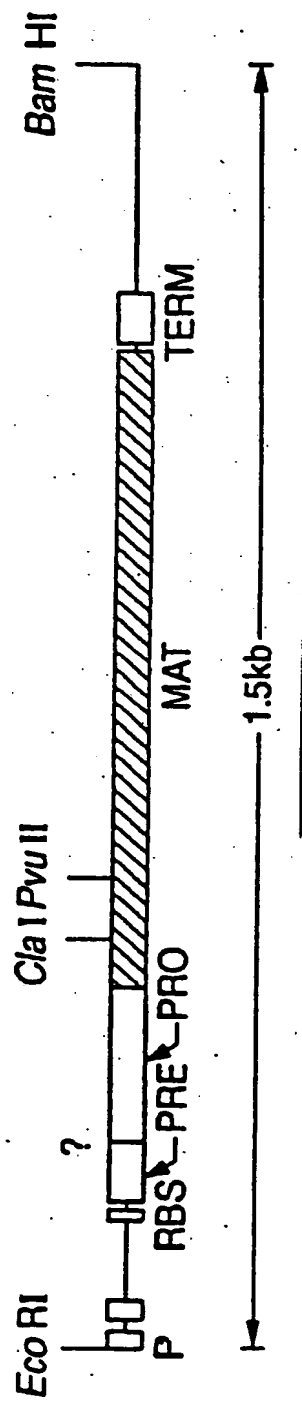


FIGURE 13.A

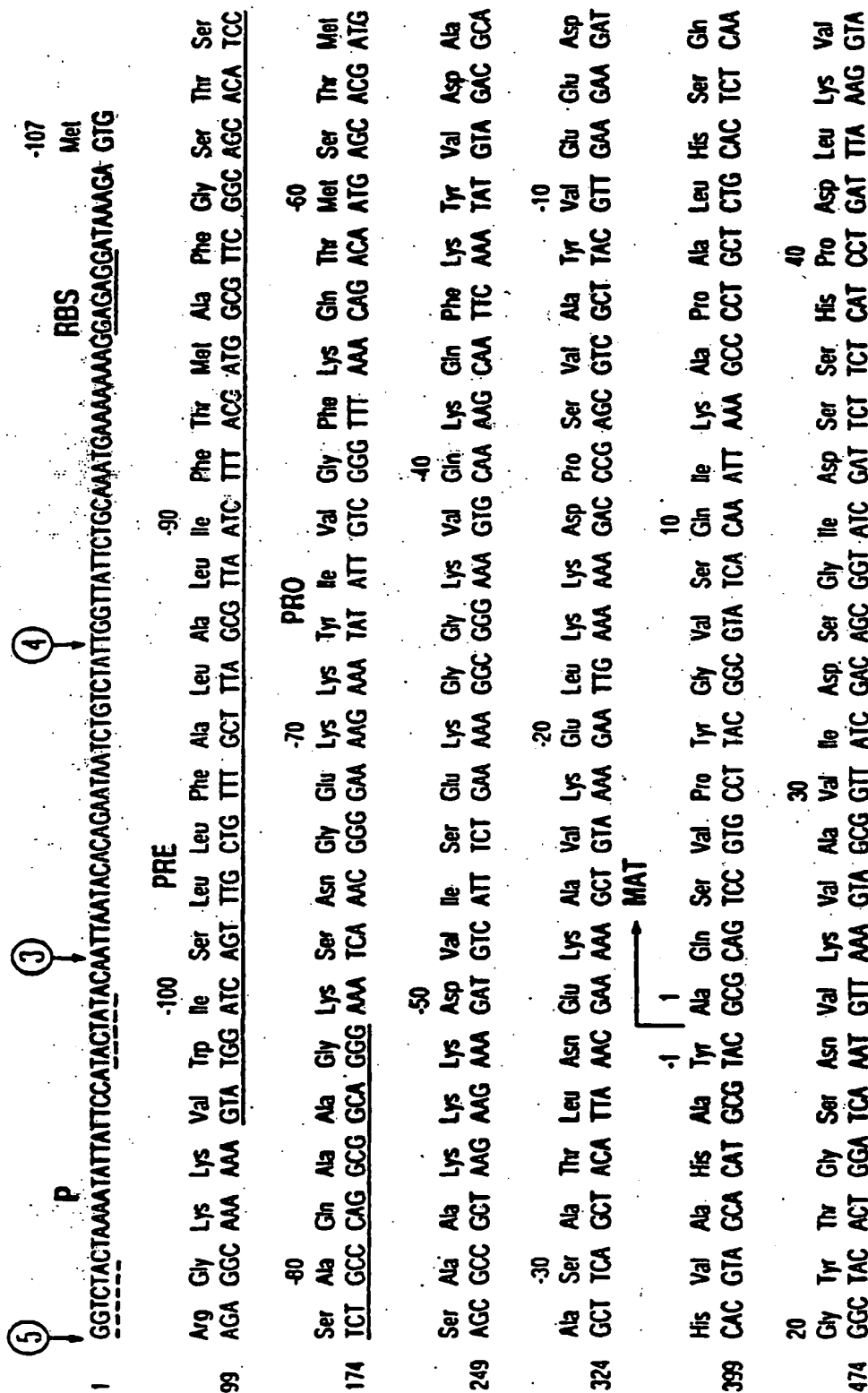


FIGURE 13.81

549 Ala Gly Gly Ala Ser Met Val Pro Ser Glu Thr Asn Pro Asn Asp 50 Asp His Gly Thr His Val Ala
 GCA GGC GGA GCC AGC ATG GTT CCT TCT GAA ACA AAT CCT TTC CAA GAC AAC AAC TCT CAC GGA ACT CAC GTT GCC

70 Gly Thr Val Ala Ala Leu Asn Asn Ser Ile Gly Val Leu Gly Val Ala Pro Ser Ala Ser Leu Tyr Ala Val Lys
 o24 GGC ACA GTT GCG GCT CTT AAT AAC TCA ATC GGT GTA TTA GGC GTT GCG CCA AGC GCA TCA CTT TAC GCT GTA AAA

100 Asp Ala 110
 Val Leu Gly Ala Asp Gly Ser Gly Gln Tyr Ser Thr Ile Ile Asn Gly Ile Glu Trp Ala Ile Ala Asn Asn Met
 699 GTT CTC GGT GCT GAC GGT TCC GGC CAA TAG AGC TGG ATC ATT AAC GGA ATC GAG TGG GCG ATC GCA AAC AAT ATG

120 Asp Val Ile Asn Met Ser Leu Gly Gly Pro Ser Gly Ser Ala Ala Leu Lys Ala Val Asp Lys Ala Val Ala
 774 GAC GTT ATT AAC ATG AGC CTC GGC GGA CCT TCT GGT TCT GCT GCT TTA AAA GCG GCA GTT GAT AAA GCC GTT GCA

150 Ser Thr 160
 Ser Gly Val Val Val Val Ala Ala Gly Asn Glu Gly Thr Ser Gly Ser Ser Thr Val Gly Tyr Pro Gly
 849 TCC GGC GTC GTA GTC GTT GCG GCA GCC GGT AAC GAA GGC ACT TCC GGC AGC TCA AGC ACA GTG GGC TAC CCT GGT

170 Lys Tyr Pro Ser Val Ile Ala Val Gly Ala Val Asp Ser Ser Asn Gln Arg Ala Ser Phe Ser Ser Val Gly Pro
 924 AAA TAC CCT TCT TCT ATT GCA GTA GGC GCT GTT GAC AGC AGC AAC CAA AGA GCA TCT TTC TCA AGC GTA GGA CCT

200 Glu Leu Asp Val Met Ala Pro Gly Val Ser Ile Gln Ser Thr Thr Leu Pro Gly Asn Lys Tyr Gly Ala Tyr Asn Gly
 999 GAG CTT GAT GTC ATG GCA CCT GGC GTA TCT ATC CAA AGC ACG CTT CCT GGA AAC AAA TAC GGG GCG TAC AAC GGT

220 Thr Ser Met Ala Ser Pro His Val Ala Gly Ala Ala Leu Ile Leu Ser Lys His Pro Asn Trp Thr Asn Thr
 1074 ACG TCA ATG GCA TCT CCG CAC GTT GCC GGA GCG GCT TTT ATT CTT TCT TCT AAG CAC CCG AAC TGG ACA AAC ACT

Figure 13.82

250 Gln 260

Gln Val Arg Ser Ser Ser Leu Glu Asn Thr Thr Thr Lys Leu Gly Asp Ser Phe Tyr Tyr Gly Lys Gly Leu Ile Asn

1149 CAA GTC CGC AGC AGT TTA GAA AAC ACC ACT ACA AAA CTT GGT GAT TCT TTC TAC TAT GGA AAA GGG CTG ATC AAC

270

Val Gln Ala Ala Ala Gln OC

1224 GTA CAG GCG GCA GCT CAG TAA AACATAAAAACGGCGCTTGGCCCCCGCGGTTTTTATTTTCTTCCCGCGAGTTCCAATCGCGTCC

1316 ATAATCGACGGATGGCTCCCTCTGAAATTTTAAACGAGAACCGGGGTTGACCCGGGCTCAGTCCCGTAACGGCCCAAGTCTGTAAACGTCTCAATCGCGCG

1416 CTCCCGGTTCCGGTCAGCTCAATGCCGTAAACGGTCGGCGGGGTTTTCTGTATACCGGGGAGACGGCATTCGTAATCGGATC

figure 13. B3

CONSERVED RESIDUES IN SUBTILISINS FROM
BACILLUS AMYLOLIQUEFACIENS

```

1      10      20
A Q S V P . G . . . . . A P A . H . . G

21     30     40
. T G S . V K V A V . D . G . . . . H P

41     50     60
D L . . . G G A S . V P . . . . . Q D

61     70     80
. N . H G T H V A G T . A A L N N S I G

81     90     100
V L G V A P S A . L Y A V K V L G A . G

101    110    120
S G . . S . L . . G . E W A . N . . . .

121    130    140
V . N . S L G . P S . S . . . . . A . .

141    150    160
. . . . . G V . V V A A . G N . G . . .

161    170    180
. . . . . Y P . . Y . . . . . A V G A .

181    190    200
D . . N . . A S F S . . G . . L D . . A

201    210    220
P G V . . Q S T . P G . . Y . . . . N G T

221    230    240
S M A . P H V A G A A A L . . . . K . . .

241    250    260
W . . . Q . R . . L . N T . . . . L G . .

261    270
. . Y G . G L . N . . A A . .

```

FIGURE 14

COMPARISON OF SUBTILISIN SEQUENCES FROM:

B. amyloliquefaciens

B. subtilis

B. licheniformis

B. lentus

01	10	20	30
1 Q S V P Y G V S Q I K A P A L H S Q G Y T G S N V K V A V I D S G I D S S H P			
A Q S V P Y G I S Q I K A P A L H S Q G Y T G S N V K V A V I D S G I D S S H P			
A Q T V P Y G I P L I K A D K V Q A Q G F K G A N V K V A V L D T G I Q A S H P			
A Q S V P W G I S R V Q A P A A H N R G L T G S G V K V A V L D T G I S T * H P			
41	50	60	70
D L K V A G G A S M V P S E T N P F Q D N N S H G T H V A G T V A A L N N S I G			
D L N V R G G A S F V P S E T N P Y Q D G S S H G T H V A G T I A A L N N S I G			
D L N V V G G A S F V A G E A Y N * T D G N G H G T H V A G T V A A L D N T T G			
D L N I R G G A S F V P G E * P S T Q D G N G H G T H V A G T I A A L N N S I G			
81	90	100	110
V L G V A P S A S L Y A V K V L G A D G S G Q Y S W I I N G I E W A I A N N M D			
V L G V S P S A S L Y A V K V L D S T G S G Q Y S W I I N G I E W A I S N N M D			
L G V A P S V S L Y A V K V L N S S G S G S Y S G I V S G I E W A T T N G M D			
V L G V A P S A E L Y A V K V L G A S G S G S V S S I A Q G L E W A G N N G M H			
121	130	140	150
V I N M S L G G P S G S A A L K A A V D K A V A S G V V V V A A A A G N E G T S G			
V I N M S L G G P T G S T A L K T V V D K A V S S G I V V V A A A A G N E G S S G			
V I N M S L G G A S G S T A M K Q A V D N A Y A R G V V V V A A A A G N S G N S G			
V A N L S L G S P S P S A T L E Q A V N S A T S R G V L V V A A S G N S G A G S			

Figure 15.A

161. 170 180 190
 S S S T V G Y P G K Y P S V I A V G A V D S S N Q R A S F S S V G P E L D V M A
 S T S T V G Y P A K Y P S T I A V G A V N S S N Q R A S F S S A G S E L D V M A
 S T N T I G Y P A K Y D S V I A V G A V D S N S N R R A S F S S V G A E L E V M A
 * * I S Y P A R Y A N A M A V G A T D Q N N R R A S F S Q Y G A G L D I V A

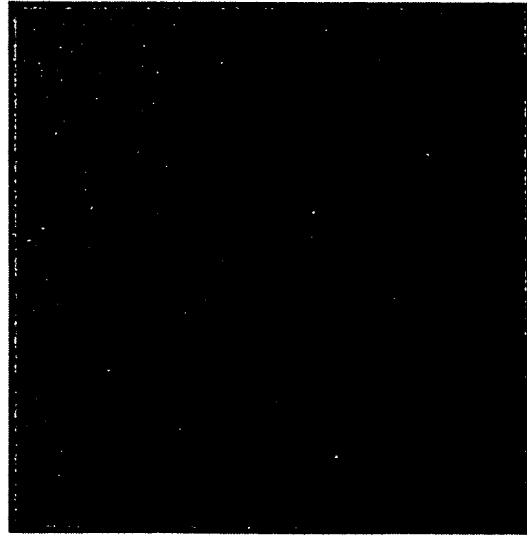
201 210 220 230
 P G V S I Q S T L P G N K Y G A Y N G T S M A S P H V A G A A A L I L S K H P N
 P G V S I Q S T L P G G T Y G A Y N G T S M A T P H V A G A A A L I L S K H P T
 P G A G V Y S T Y P P T N T Y A T L N G T S M A S P H V A G A A A L I L S K H P N
 P G V N V Q S T Y P G S T Y A S L N G T S M A T P H V A G A A A L V K Q K N P S

241 250 260 270
 W T N T Q V R S S L E N T T K L G D S F Y Y G K G L I N V Q A A A Q
 W T N A Q V R D R L E S T A T Y L G N S F Y Y G K G L I N V Q A A A Q
 L S A S Q V R N R L S S T A T Y L G S S F Y Y G K G L I N V E A A A Q
 W S N V Q I R N N H L K N T A T S L G S T N L Y G S G L V N A E A A T R

Figure 15.B

Initial evaluation results

MC-A



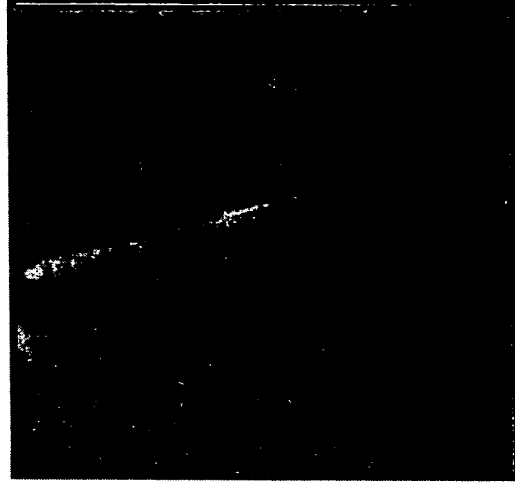
m mbh 2 4 6 8 10 12 P m

MC-3



m mbh 2 4 6 8 10 12 P m

MC-4

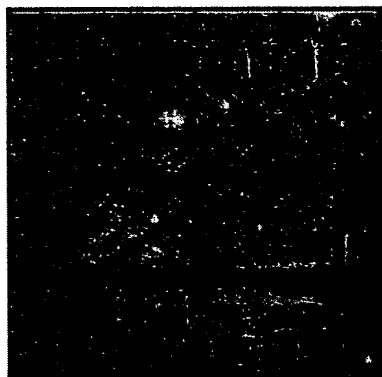


m mbh 2 4 6 8 10 12 P m

Fig. 16

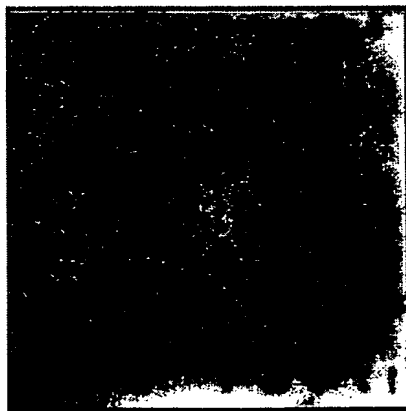
Comparison with Properase

Properase 60°C 30 minutes



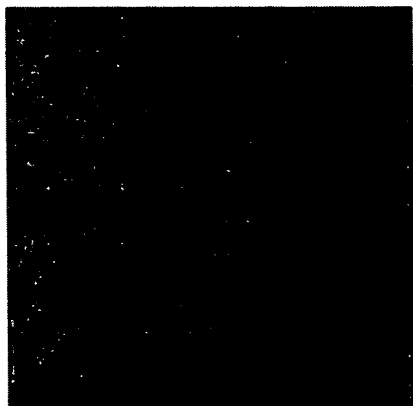
m 2 4 6 8 10 12 P rPrP m

MC-A 50°C 30 minutes



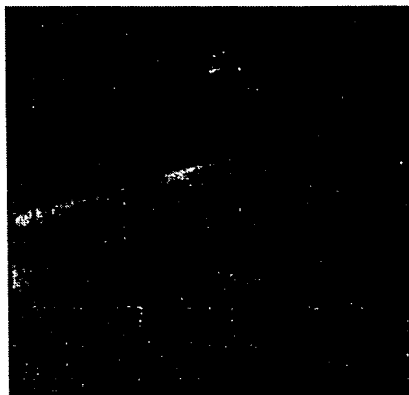
m 2 4 6 8 10 12 P rPrP m

MC-3 50°C 30 minutes



m 2 4 6 8 10 12 P rPrP m

MC-4 50°C 30 minutes



m mbh 2 4 6 8 10 12 P m

Fig. 17

Comparison with Properase

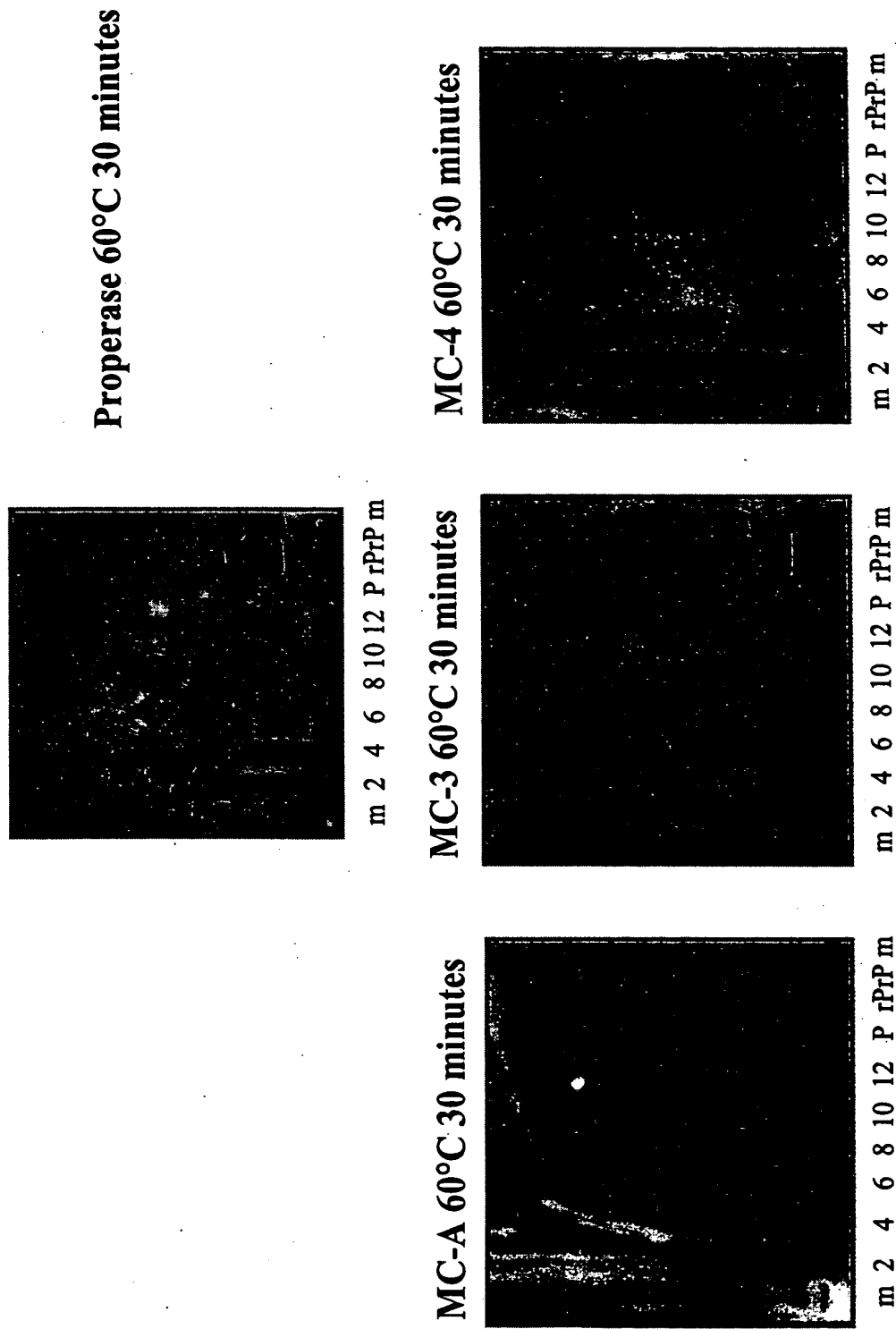
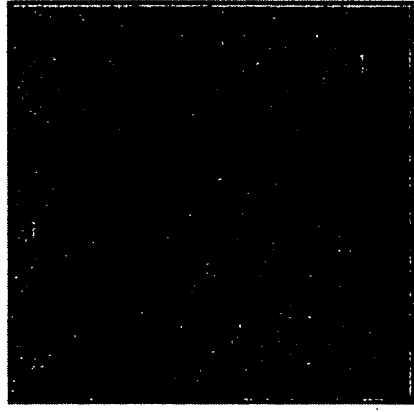


Fig. 18

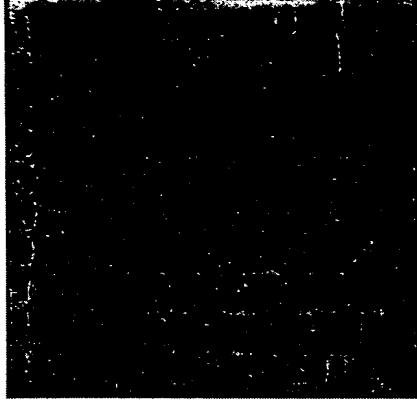
Temperature profiling with MC-3

50°C 30 minutes



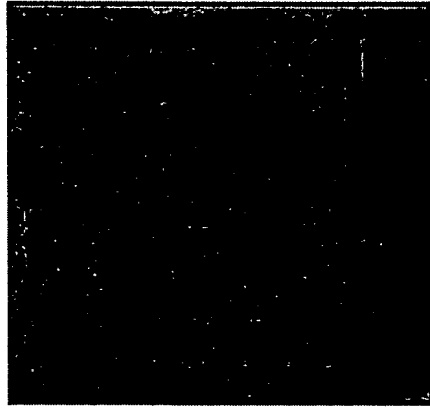
m 2 4 6 8 10 12 P rPrP m

70°C 30 minutes



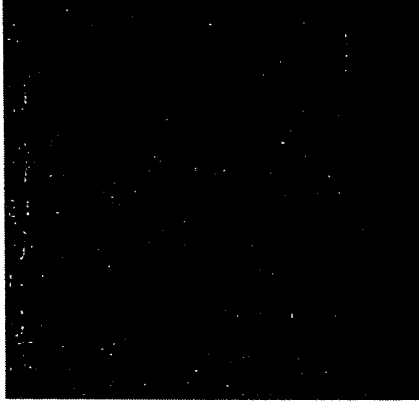
m 2 4 6 8 10 12 P rPrP m

60°C 30 minutes



m 2 4 6 8 10 12 P rPrP m

80°C 30 minutes



m 2 4 6 8 10 12 P rPrP m

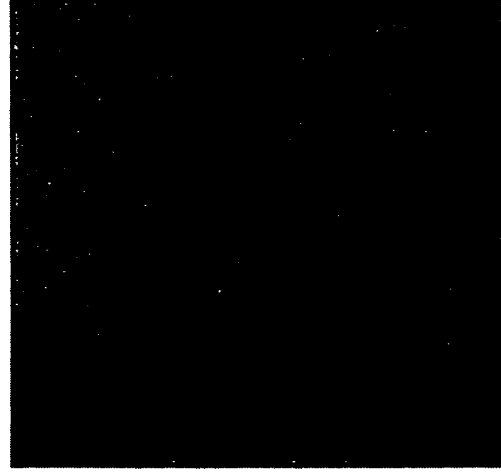
Fig. 19

Detection with PAb2

mbh pH 2-12 digested at 50°C 30 minutes

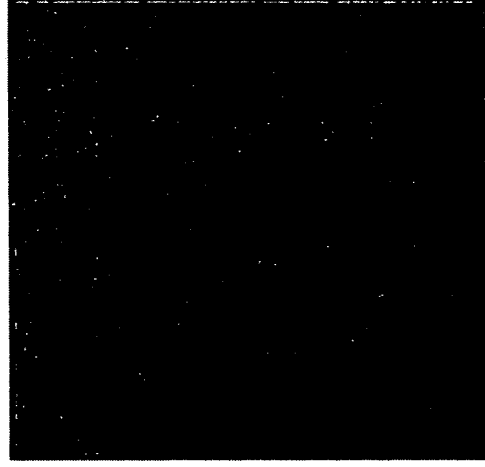
- Detected with a chemiluminescent detection substrate (Pierce)

MC-A



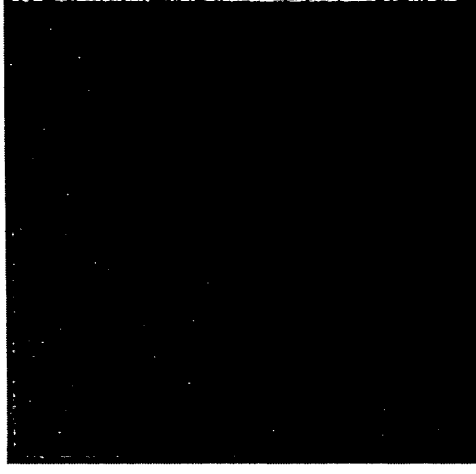
m 2 4 6 8 10 12 P rPrP m

MC-3



m 2 4 6 8 10 12 P rPrP

MC-4

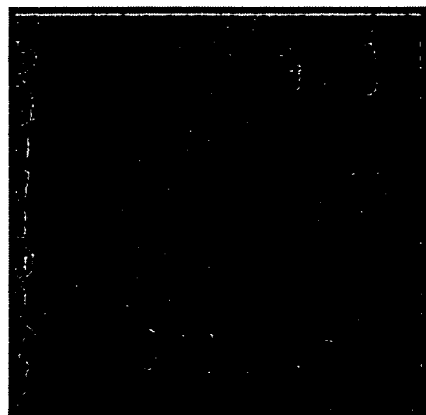


m mbh 2 4 6 8 10 12 P m

Fig. 20

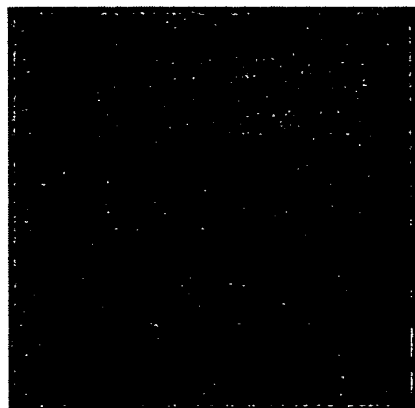
MC-3 dilutions at pH 10 & pH 12

6H4 West Dura



m
rPrP
P
1:100
1:50
1:20
1:4
n
mbh
m

PAb2 West Dura



m
rPrP
P
1:100
1:50
1:20
1:4
n
mbh
m

pH 10

Monomer bands at
1:20 dilution

HMW bands across
dilution range

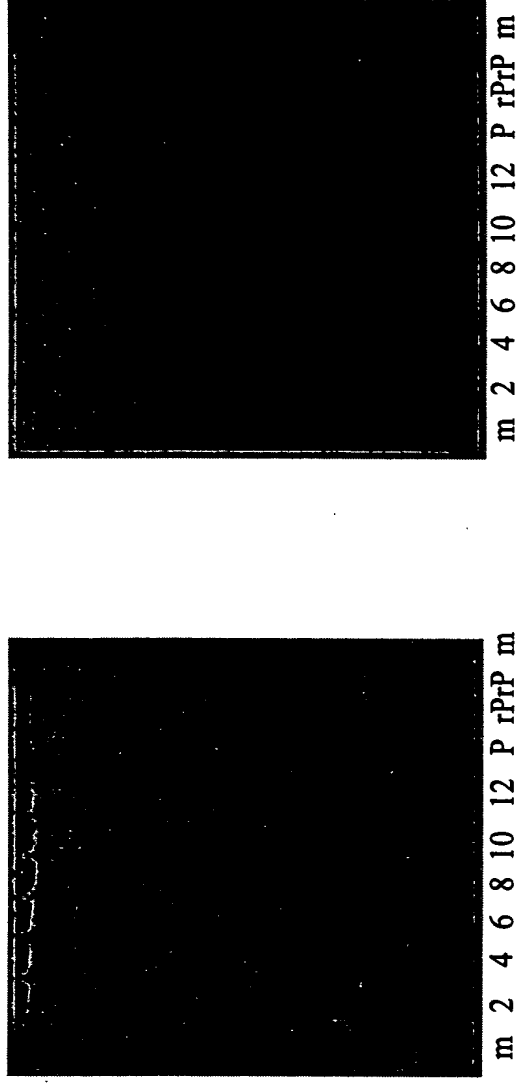
pH 12

No monomer bands

HMW bands much
reduced across
dilution range

Fig. 21

Comparison with Proteinase K



- Characteristic PrP^{Sc} monomer bands pH 2-10
Incomplete digestion pH12 however no clear monomers
HMW bands present pH 2-12
The new proteases are better at removing both the
monomer and HMW bands than Proteinase K

Fig. 22